

# House Concurrent Resolution No. 48

## 95TH GENERAL ASSEMBLY

INTRODUCED BY REPRESENTATIVES PARSON (Sponsor), TALBOY, STILL, RUCKER, SILVEY,  
KANDER, SCHOELLER, JONES (117), POLLOCK, ICET, BRUNS,  
HOLSMAN AND WEBBER (Co-sponsors).

2490L.011

2 **Whereas**, it is the goal of the Missouri General Assembly to promote better health  
3 outcomes for Missourians by exploring innovation and life sciences, high-tech jobs, and  
4 economic growth; and

5 **Whereas**, bioscience firms conducting business in Missouri are actively facilitating  
6 life science discovery to develop medicines targeting cancers, rare diseases, and other critical  
7 conditions which will lead to significant economic development opportunities; and

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9 **Whereas**, biopharmaceutical companies supported a total of 65,192 jobs in  
10 Missouri in 2006 - 10,884 directly in the sector and 54,307 in other sectors; and

11  
12 **Whereas**, direct biopharmaceutical wages in Missouri were estimated to be \$1  
13 billion in 2006, resulting in an estimated \$251.1 million in federal taxes and \$39.5 million in  
14 state taxes; and

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16 **Whereas**, in 2008, United States scientists and researchers were conducting 21,795  
17 studies to develop medicines targeting cancers, rare diseases, and other important conditions,  
18 with 2,397 of these trials active in Missouri; and

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20 **Whereas**, Missouri has research universities and bioscience companies which play  
21 a key role in the discovery of cures and treatments for diseases that were previously thought  
22 incurable, and the Missouri General Assembly wishes to ensure that these universities and  
23 companies remain viable to develop medicines that promote public health, encourage job  
24 creation, and foster economic growth; and

25       **Whereas**, on average, it requires over \$1.2 billion and 10-15 years to discover,  
26 develop, and secure United States Food and Drug Administration (FDA) approval to bring a new  
27 biologic medicine to the market, and it is imperative that bioscience companies are able to  
28 recoup sufficient financial returns to justify investing in this critically important enterprise; and  
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30       **Whereas**, biologics are today's most advanced medicines and include many of the  
31 latest breakthrough medical therapies for serious and life-threatening illnesses such as cancer,  
32 multiple sclerosis, diabetes, HIV/AIDS, and many serious rare diseases; and  
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34       **Whereas**, the Missouri General Assembly believes that balanced biosimilar  
35 legislation should include a provision that provides pioneer bioscience companies with 14 years  
36 of exclusive use of the costly, proprietary data they must produce to secure FDA approval before  
37 a biosimilar manufacturer may acquire this data without cost; and  
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39       **Whereas**, the Missouri General Assembly wishes to ensure that patient safety is  
40 the primary concern of the FDA when approving both pioneer and biosimilar drugs, and the  
41 Missouri General Assembly recognizes that biosimilar manufacturers may not be required to  
42 duplicate all tests required of the pioneer manufacturer; and  
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44       **Whereas**, the Missouri General Assembly believes that all Missourians will benefit  
45 from more affordable and more innovative biological medicines to improve health outcomes, as  
46 well as by the creation of jobs and economic growth associated with bioscience research and  
47 development activities conducted by universities and bioscience industry:  
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49       **Now, therefore, be it resolved** that the members of the House of  
50 Representatives of the Ninety-fifth General Assembly, First Regular Session, the Senate  
51 concurring therein, hereby urges the United States Congress to adopt balanced biosimilar  
52 legislation that provides reasonable incentives that will foster the research and development of  
53 next generation, life-saving biological medicines as well as job creation and economic  
54 expansion, and encourages the creation of a transparent, science-based regulatory review system  
55 that will allow a fair and prompt FDA review of biosimilar products so consumers may benefit  
56 from increased price competition as soon as appropriate; and

57       **Be it further resolved** that the Chief Clerk of the Missouri House of  
58 Representatives be instructed to prepare properly inscribed copies of this resolution for the  
59 President of the United States Senate, the Speaker of the United States House of Representatives,  
60 and each member of the Missouri Congressional delegation.

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